FDA 510(k) Premarket Notification

K222780 LILY Extension Tube and Needleless Connector

CLEARED

Nov 9, 2023
421 days to decision

K222780 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k222780/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 14, 2022
Decision date	Nov 9, 2023
Days to decision	421 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Lily Medical Corporation**
Location	Miaoli County, CN
Contact	Steven Shen
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	**Benq Medical Technology Corporation**
Contact	Steven Shen

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov